## AMENDMENTS TO THE CLAIMS

This listing replaces all prior versions and listings of claims in the application.

## **Listing of Claims**

## 1. (Currently amended) A compound of formula I

$$R^1$$
  $NR^3R^4$   $I$ 

or a pharmaceutically acceptable salt, hydrate, solvate or prodrug of the compound, wherein:

 $R^1$  is hydrogen, -OH, -NO<sub>2</sub>, -CN, -COOR, -OCH<sub>2</sub>OR,  $C_1$ - $C_6$  alkyl,  $C_2$ - $C_6$  alkenyl,  $C_2$ - $C_6$  alkynyl,  $C_1$ - $C_6$  alkoxy or halo;

R is  $C_1$ - $C_6$  alkyl;

R<sup>2</sup> is a non-radioactive halo or a radioactive halo;

R<sup>3</sup> is hydrogen, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl or C<sub>2</sub>-C<sub>6</sub> alkynyl; and

R<sup>4</sup> is hydrogen, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl or C<sub>2</sub>-C<sub>6</sub> alkynyl, wherein the alkyl, alkenyl or alkynyl comprises a radioactive carbon or is substituted with a radioactive halo when R<sup>2</sup> is hydrogen or a non-radioactive halo;

provided that when R<sup>1</sup> is hydrogen or -OH, R<sup>2</sup> is hydrogen and R<sup>4</sup> is -<sup>11</sup>CH<sub>3</sub>, then R<sup>3</sup> is C<sub>2</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl or C<sub>2</sub>-C<sub>6</sub> alkynyl; and

further provided that when R<sup>1</sup> is hydrogen, R<sup>2</sup> hydrogen and R<sup>4</sup> is -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub><sup>18</sup>F, then R<sup>3</sup> is C<sub>2</sub>-C<sub>6</sub>-alkyl, C<sub>2</sub>-C<sub>6</sub>-alkenyl or C<sub>2</sub>-C<sub>6</sub>-alkynyl.

2. (Withdrawn) The compound of claim 1, wherein:

R<sup>1</sup> is hydrogen, -OH, -CN, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>1</sub>-C<sub>6</sub> alkoxy or halo; and R<sup>4</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl or C<sub>2</sub>-C<sub>6</sub> alkynyl, wherein the alkyl, alkenyl or alkynyl comprises a radioactive carbon.

3. (Withdrawn) The compound of claim 2, wherein:

$$R^1$$
 is hydrogen, -OH, -CN, -OCH<sub>3</sub>, -CH<sub>3</sub> or -Br; and  $R^3$  is hydrogen or -CH<sub>3</sub>; and  $R^4$  is  $-^{11}$ CH<sub>3</sub>.

4. (Original) The compound of claim 1, wherein:

R<sup>2</sup> is a non-radioactive halo or a radioactive halo, wherein the halo is iodo; and

 $R^4$  is hydrogen,  $C_1$ - $C_6$  alkyl,  $C_2$ - $C_6$  alkenyl or  $C_2$ - $C_6$  alkynyl, wherein the alkyl, alkenyl or alkynyl comprises a radioactive carbon when  $R^2$  is a non-radioactive halo.

5. (Withdrawn) The compound of claim 4, wherein:

6. (Withdrawn) The compound of claim 5, wherein:

$$R^1$$
 is -OH or  $C_1$ - $C_6$  alkoxy; 
$$R^2$$
 is a radioiodine; and 
$$R^3$$
 and  $R^4$  are independently hydrogen or  $C_1$ - $C_6$  alkyl.

7. (Withdrawn) The compound of claim 6, wherein:

$$R^1$$
 is  $-OH$ ;

$$R^2$$
 is  $-^{123}I$  or  $-^{125}I$ ; and

R<sup>3</sup> and R<sup>4</sup> are each hydrogen.

- 8. (Original) The compound of claim 1, wherein R<sup>2</sup> is a radiofluoro.
- 9. (Original) The compound of claim 8, wherein:

$$R^1$$
 is –OH or  $C_1$ - $C_6$  alkoxy;

R<sup>3</sup> and R<sup>4</sup> are independently hydrogen or C<sub>1</sub>-C<sub>6</sub> alkyl.

10. (Withdrawn) The compound of claim 9, wherein:

$$R^1$$
 is  $-OH$ ;

R<sup>3</sup> is hydrogen; and

$$R^4$$
 is  $-CH_3$ .

- 11. (Withdrawn) The compound of claim 1, wherein  $R^4$  is  $C_1$ - $C_6$  alkyl,  $C_2$ - $C_6$  alkenyl or  $C_2$ - $C_6$  alkynyl, wherein the alkyl, alkenyl or alkynyl is substituted with a radioactive halo.
- 12. (Withdrawn) The compound of claim 11, wherein:

R<sup>2</sup> is hydrogen;

R<sup>3</sup> is hydrogen or C<sub>1</sub>-C<sub>6</sub> alkyl; and

R<sup>4</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl substituted with <sup>18</sup>F.

13. (Withdrawn) The compound of claim 12, wherein:

$$R^1$$
 is  $-OH$ ;

R<sup>3</sup> is hydrogen; and

R<sup>4</sup> is -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub><sup>18</sup>F.

- 14. (Original) A pharmaceutical composition comprising
  - (i) an effective amount of a compound of claim 1; and
  - (ii) a pharmaceutically acceptable carrier.
- 15. (Withdrawn) A method for detecting amyloid deposit(s) in vivo, comprising:
  - (i) administering to a mammal an effective amount of a compound of claim 1, wherein the compound would bind any amyloid deposit(s) in the mammal; and
  - (ii) detecting binding of the compound to amyloid deposit(s) in the mammal.
- 16. (Withdrawn) The method of claim 15, wherein the amyloid deposit(s) is/are located in the brain of the mammal.
- 17. (Withdrawn) The method of claim 15, wherein the mammal is a human who is suspected of having Alzheimer's disease, familial Alzheimer's disease, Down's syndrome, Mild Cognitive Impairment or homozygotes for apolipoprotein E4 allele.
- 18. (Withdrawn) The method of claim 15, wherein the detecting is effected by gamma imaging, magnetic resonance imaging or magnetic resonance spectroscopy.
- 19. (Withdrawn) The method of claim 18, wherein the detecting is effected by gamma imaging.
- 20. (Withdrawn) The method of claim 19, wherein the gamma imaging is PET or SPECT.
- 21. (Withdrawn) The method of claim 15, wherein the compound is administered intravenously.
- 22. (Withdrawn) A method for detecting amyloid deposit(s) in vitro comprising:
  - (i) contacting a bodily tissue with an effective amount of a compound of claim 1, wherein the compound would bind any amyloid deposit(s) in the tissue; and
  - (ii) detecting binding of the compound to amyloid deposit(s) in the tissue.

- 23. (Withdrawn) The method of claim 22, wherein the compound is in a solution that further comprises 25-99% ethanol, with the remainder of the solution being water.
- 24. (Withdrawn) The method of claim 23, wherein the solution comprises 0-50% ethanol and 0.0001 to  $100 \,\mu\text{M}$  of the compound.
- 25. (Withdrawn) The method of claim 22 wherein the detecting is effected by bright-field, fluorescence, laser-confocal or cross-polarization microscopy.
- 26. (Withdrawn) The method of claim 22, wherein the method further comprises:
  - (iii) separating from the tissue the amyloid deposit(s) bound to the compound; and
  - (iv) quantifying the amyloid deposit(s) bound to the compound.
- 27. (Withdrawn) A method for distinguishing an Alzheimer's diseased brain from a normal brain comprising:
  - (i) obtaining tissues from (i) the cerebellum and (ii) another area of the same brain, of a normal mammal and of a mammal suspected of having Alzheimer's disease;
  - (ii) contacting the tissues with a compound of claim 1;
  - (iii) quantifying the amyloid bound to the compound;
  - (iv) calculating the ratio of (a) the amount of amyloid in the area of the brain other than the cerebellum to (b) the amount of amyloid in the cerebellum;
  - (v) comparing the ratio for a normal mammal with the ratio for a mammal suspected of having Alzheimer's disease.